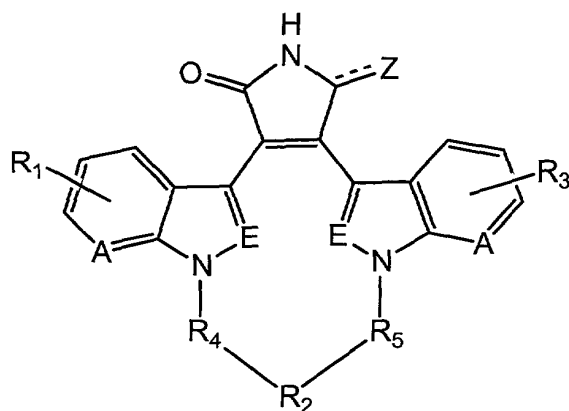


What is Claimed is:

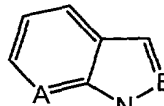
1. A compound of Formula (I):



Formula (I)

wherein

A and E are independently selected from the group consisting of a hydrogen substituted

- 5 carbon atom and a nitrogen atom; wherein  is independently selected from the group consisting of 1*H*-indole, 1*H*-pyrrolo[2,3-*b*]pyridine, 1*H*-pyrazolo[3,4-*b*]pyridine and 1*H*-indazole;

Z is selected from O or dihydro; wherein when Z is dihydro, each hydrogen atom is attached by a single bond;

- 10 R_4 and R_5 are independently selected from C_{1-8} alkyl, C_{2-8} alkenyl and C_{2-8} alkynyl, wherein R_4 and R_5 are optionally substituted with oxo;

- 15 R_2 is selected from the group consisting of $-C_{1-8}$ alkyl-, $-C_{2-8}$ alkenyl-, $-C_{2-8}$ alkynyl-, $-O-(C_{1-8})alkyl-O-$, $-O-(C_{2-8})alkenyl-O-$, $-O-(C_{2-8})alkynyl-O-$, $-C(O)-(C_{1-8})alkyl-C(O)-$ (wherein any of the foregoing alkyl, alkenyl and alkynyl linking groups are straight carbon chains optionally substituted with one to four substituents independently selected from the group consisting of C_{1-8} alkyl, C_{1-8} alkoxy, C_{1-8} alkoxy(C_{1-8})alkyl, carboxyl, carboxyl(C_{1-8})alkyl, $-C(O)O-(C_{1-8})alkyl$, $-C_{1-8}alkyl-C(O)O-(C_{1-8})alkyl$, amino (substituted with a substituent independently selected from the group consisting of hydrogen and

C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy, hydroxy(C₁₋₈)alkyl and oxo; and, wherein any of the foregoing alkyl, alkenyl and alkynyl linking groups are optionally substituted with one to two substituents independently selected from the group consisting of heterocyclyl, aryl, heteroaryl, heterocyclyl(C₁₋₈)alkyl, aryl(C₁₋₈)alkyl, heteroaryl(C₁₋₈)alkyl, spirocycloalkyl and spiroheterocyclyl (wherein any of the foregoing cycloalkyl, heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein any of the foregoing heterocyclyl substituents are optionally substituted with oxo)), cycloalkyl, heterocyclyl, aryl, heteroaryl (wherein cycloalkyl, heterocyclyl, aryl and heteroaryl are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein heterocyclyl is optionally substituted with oxo), -(O-(CH₂)₁₋₆)₀₋₅-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -(O-(CH₂)₁₋₆)₀₋₅-NR₆-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-NR₆-, -(O-(CH₂)₁₋₆)₀₋₅-S-, -O-(CH₂)₁₋₆-S-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-S-, -NR₆-, -NR₆-NR₇-, -NR₆-(CH₂)₁₋₆-NR₇-, -NR₆-(CH₂)₁₋₆-NR₇-(CH₂)₁₋₆-NR₈-, -NR₆-C(O)-, -C(O)-NR₆-, -C(O)-(CH₂)₀₋₆-NR₆-(CH₂)₀₋₆-C(O)-, -NR₆-(CH₂)₀₋₆-C(O)-(CH₂)₁₋₆-C(O)-(CH₂)₀₋₆-NR₇-, -NR₆-C(O)-NR₇-, -NR₆-C(NR₇)-NR₈-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S-, -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S-, -NR₆-(CH₂)₁₋₆-S-(CH₂)₁₋₆-NR₇- and -SO₂- (wherein

R₆, R₇ and R₈ are independently selected from the group consisting of hydrogen, C₁₋₈alkyl, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl(C₁₋₈)alkyl, amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), hydroxy(C₁₋₈)alkyl, heterocyclyl(C₁₋₈)alkyl, aryl(C₁₋₈)alkyl and heteroaryl(C₁₋₈)alkyl (wherein the foregoing heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein heterocyclyl is optionally substituted with oxo));

with the proviso that, if A and E are selected from a hydrogen substituted carbon atom, then R₂ is selected from the group consisting of -C₂₋₈alkynyl-, -O-(C₁₋₈)alkyl-O-, -O-(C₂₋₈)alkenyl-O-, -O-(C₂₋₈)alkynyl-O-, -C(O)-(C₁₋₈)alkyl-C(O)- (wherein any of the foregoing alkyl, alkenyl and alkynyl linking groups are straight carbon chains optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, -C(O)O-(C₁₋₈)alkyl, -C₁₋₈alkyl-C(O)O-(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy, hydroxy(C₁₋₈)alkyl and oxo; and, wherein any of the foregoing alkyl, alkenyl and alkynyl linking groups are optionally substituted with one to two substituents independently selected from the group consisting of heterocyclyl, aryl, heteroaryl, heterocyclyl(C₁₋₈)alkyl, aryl(C₁₋₈)alkyl, heteroaryl(C₁₋₈)alkyl, spirocycloalkyl and spiroheterocyclyl (wherein any of the foregoing cycloalkyl, heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and

C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein any of the foregoing heterocyclyl substituents are optionally substituted with oxo)), cycloalkyl (wherein cycloalkyl is optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl), -(O-(CH₂)₁₋₆)₁₋₅-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -(O-(CH₂)₁₋₆)₁₋₅-NR₆-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-NR₆-, -(O-(CH₂)₁₋₆)₀₋₅-S-, -O-(CH₂)₁₋₆-S-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-S-, -NR₆-NR₇-, -NR₆-(CH₂)₁₋₆-NR₇-, -NR₆-(CH₂)₁₋₆-NR₇-(CH₂)₁₋₆-NR₈-, -NR₉-C(O)-, -C(O)-NR₉-, -C(O)-(CH₂)₀₋₆-NR₆-(CH₂)₀₋₆-C(O)-, -NR₆-(CH₂)₀₋₆-C(O)-(CH₂)₁₋₆-C(O)-(CH₂)₀₋₆-NR₇-, -NR₆-C(O)-NR₇-, -NR₆-C(NR₇)-NR₈-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S-, -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S- and -NR₆-(CH₂)₁₋₆-S-(CH₂)₁₋₆-NR₇- (wherein R₆, R₇ and R₈ are independently selected from the group consisting of hydrogen, C₁₋₈alkyl, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl(C₁₋₈)alkyl, amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), hydroxy(C₁₋₈)alkyl, heterocyclyl(C₁₋₈)alkyl, aryl(C₁₋₈)alkyl and heteroaryl(C₁₋₈)alkyl (wherein the foregoing heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein heterocyclyl is optionally substituted with oxo); and, wherein R₉ is selected from the group

consisting of C₁₋₈alkyl, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl(C₁₋₈)alkyl, amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), hydroxy(C₁₋₈)alkyl, heterocyclyl(C₁₋₈)alkyl, aryl(C₁₋₈)alkyl and heteroaryl(C₁₋₈)alkyl (wherein the foregoing heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein heterocyclyl is optionally substituted with oxo)); and,

R₁ and R₃ are independently selected from the group consisting of hydrogen, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl (wherein alkyl, alkenyl and alkynyl are optionally substituted with a substituent selected from the group consisting of C₁₋₈alkoxy, alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), (halo)₁₋₃, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy, hydroxy(C₁₋₈)alkyl and oxo), C₁₋₈alkoxy, C₁₋₈alkoxycarbonyl, (halo)₁₋₃(C₁₋₈)alkoxy, C₁₋₈alkylthio, aryl, heteroaryl (wherein aryl and heteroaryl are optionally substituted with a substituent selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl), amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), cyano, halogen, hydroxy and nitro;

and pharmaceutically acceptable salts thereof.

2. The compound of claim 1 wherein R₄ and R₅ are independently selected from C₁₋₆alkyl, C₂₋₆alkenyl and C₂₋₆alkynyl optionally substituted with oxo.
3. The compound of claim 1 wherein R₄ and R₅ are independently selected from C₁₋₆alkyl, C₂₋₆alkenyl and C₂₋₆alkynyl.
4. The compound of claim 1 wherein R₄ and R₅ are independently selected from C₁₋₆alkyl.
5. The compound of claim 1 wherein R₂ is selected from the group consisting of -C₁₋₈alkyl-, -C₂₋₄alkenyl-, -C₂₋₄alkynyl-, -O-(C₁₋₄)alkyl-O-, -O-(C₂₋₄)alkenyl-O-, -O-(C₂₋₄)alkynyl-O-, -C(O)-(C₁₋₄)alkyl-C(O)- (wherein any of the foregoing alkyl, alkenyl and alkynyl linking groups are straight carbon chains optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkoxy(C₁₋₄)alkyl, carboxyl, carboxyl(C₁₋₄)alkyl, -C(O)O-(C₁₋₄)alkyl, -C₁₋₄alkyl-C(O)O-(C₁₋₄)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₄)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₄)alkyl, (halo)₁₋₃(C₁₋₄)alkoxy, hydroxy, hydroxy(C₁₋₄)alkyl and oxo; and, wherein any of the foregoing alkyl, alkenyl and alkynyl linking groups are optionally substituted with one to two substituents independently selected from the group consisting of heterocyclyl, aryl, heteroaryl, heterocyclyl(C₁₋₄)alkyl, aryl(C₁₋₄)alkyl, heteroaryl(C₁₋₄)alkyl, spirocycloalkyl and spiroheterocyclyl (wherein any of the foregoing cycloalkyl, heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkoxy(C₁₋₄)alkyl, carboxyl, carboxyl(C₁₋₄)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₄)alkyl (wherein amino is

substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₄)alkyl, (halo)₁₋₃(C₁₋₄)alkoxy, hydroxy and hydroxy(C₁₋₄)alkyl; and, wherein any of the foregoing heterocyclyl substituents are optionally substituted with oxo)), cycloalkyl, heterocyclyl, aryl, heteroaryl (wherein cycloalkyl, heterocyclyl, aryl and heteroaryl are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkoxy(C₁₋₄)alkyl, carboxyl, carboxyl(C₁₋₄)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₄)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₄)alkyl, (halo)₁₋₃(C₁₋₄)alkoxy, hydroxy and hydroxy(C₁₋₄)alkyl; and, wherein heterocyclyl is optionally substituted with oxo), -(O-(CH₂)₁₋₆)₀₋₅-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆)₀₋₅-NR₆-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-NR₆-, -(O-(CH₂)₁₋₆)₀₋₅-S-, -O-(CH₂)₁₋₆-S-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-S-, -NR₆-, -NR₆-NR₇-, -NR₆-(CH₂)₁₋₆-NR₇-, -NR₆-(CH₂)₁₋₆-NR₇-(CH₂)₁₋₆-NR₈-, -NR₆-C(O)-, -C(O)-NR₆-, -C(O)-(CH₂)₀₋₆-NR₆-(CH₂)₀₋₆-C(O)-, -NR₆-(CH₂)₀₋₆-C(O)-(CH₂)₁₋₆-C(O)-(CH₂)₀₋₆-NR₇-, -NR₆-C(O)-NR₇-, -NR₆-C(NR₇)-NR₈-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S-, -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S-, -NR₆-(CH₂)₁₋₆-S-(CH₂)₁₋₆-NR₇- and -SO₂- (wherein R₆, R₇ and R₈ are independently selected from the group consisting of hydrogen, C₁₋₄alkyl, C₁₋₄alkoxy(C₁₋₄)alkyl, carboxyl(C₁₋₄)alkyl, amino(C₁₋₄)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), hydroxy(C₁₋₄)alkyl, heterocyclyl(C₁₋₄)alkyl, aryl(C₁₋₄)alkyl and heteroaryl(C₁₋₄)alkyl (wherein the foregoing heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkoxy(C₁₋₄)alkyl, carboxyl, carboxyl(C₁₋₄)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₄)alkyl (wherein amino is substituted with a substituent

independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₄)alkyl, (halo)₁₋₃(C₁₋₄)alkoxy, hydroxy and hydroxy(C₁₋₄)alkyl; and, wherein heterocyclyl is optionally substituted with oxo));

- 5 with the proviso that, if A and E are selected from a hydrogen substituted carbon atom, then R₂ is selected from the group consisting of -C₂₋₄alkynyl-, -O-(C₁₋₄)alkyl-O-, -O-(C₂₋₄)alkenyl-O-, -O-(C₂₋₄)alkynyl-O-, -C(O)-(C₁₋₄)alkyl-C(O)- (wherein any of the foregoing alkyl, alkenyl and alkynyl linking groups are straight carbon chains optionally substituted with one
- 10 to four substituents independently selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkoxy(C₁₋₄)alkyl, carboxyl, carboxyl(C₁₋₄)alkyl, -C(O)O-(C₁₋₄)alkyl, -C₁₋₄alkyl-C(O)O-(C₁₋₄)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₄)alkyl (wherein amino is substituted with a substituent
- 15 independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₄)alkyl, (halo)₁₋₃(C₁₋₄)alkoxy, hydroxy, hydroxy(C₁₋₄)alkyl and oxo; and, wherein any of the foregoing alkyl, alkenyl and alkynyl linking groups are optionally substituted with one to two substituents independently selected from the group consisting of heterocyclyl, aryl, heteroaryl,
- 20 heterocyclyl(C₁₋₄)alkyl, aryl(C₁₋₄)alkyl, heteroaryl(C₁₋₄)alkyl, spirocycloalkyl and spiroheterocyclyl (wherein any of the foregoing cycloalkyl, heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkoxy(C₁₋₄)alkyl, carboxyl, carboxyl(C₁₋₄)alkyl, amino
- 25 (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₄)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₄)alkyl, (halo)₁₋₃(C₁₋₄)alkoxy, hydroxy and hydroxy(C₁₋₄)alkyl; and, wherein any of the foregoing heterocyclyl substituents
- 30 are optionally substituted with oxo)), cycloalkyl (wherein cycloalkyl is optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkoxy(C₁₋₄)alkyl, carboxyl, carboxyl(C₁₋₄)alkyl, amino (substituted with a substituent independently

selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₄)alkyl
 (wherein amino is substituted with a substituent independently selected from the
 group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₄)alkyl,
 (halo)₁₋₃(C₁₋₄)alkoxy, hydroxy and hydroxy(C₁₋₄)alkyl), -(O-(CH₂)₁₋₆)₁₋₅-O-,
 5 -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-,
 -(O-(CH₂)₁₋₆)₁₋₅-NR₆-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-,
 -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-NR₆-, -(O-(CH₂)₁₋₆)₀₋₅-S-, -O-(CH₂)₁₋₆-S-(CH₂)₁₋₆-O-,
 -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-S-, -NR₆-NR₇-, -NR₆-(CH₂)₁₋₆-NR₇-,
 -NR₆-(CH₂)₁₋₆-NR₇-(CH₂)₁₋₆-NR₈-, -NR₉-C(O)-, -C(O)-NR₉-,
 10 -C(O)-(CH₂)₀₋₆-NR₆-(CH₂)₀₋₆-C(O)-,
 -NR₆-(CH₂)₀₋₆-C(O)-(CH₂)₁₋₆-C(O)-(CH₂)₀₋₆-NR₇-, -NR₆-C(O)-NR₇-,
 -NR₆-C(NR₇)-NR₈-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S-, -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-,
 -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S- and -NR₆-(CH₂)₁₋₆-S-(CH₂)₁₋₆-NR₇- (wherein R₆,
 R₇ and R₈ are independently selected from the group consisting of hydrogen,
 15 C₁₋₄alkyl, C₁₋₄alkoxy(C₁₋₄)alkyl, carboxyl(C₁₋₄)alkyl, amino(C₁₋₄)alkyl (wherein
 amino is substituted with a substituent independently selected from the group
 consisting of hydrogen and C₁₋₄alkyl), hydroxy(C₁₋₄)alkyl,
 heterocyclyl(C₁₋₄)alkyl, aryl(C₁₋₄)alkyl and heteroaryl(C₁₋₄)alkyl (wherein the
 foregoing heterocyclyl, aryl and heteroaryl substituents are optionally
 20 substituted with one to four substituents independently selected from the group
 consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkoxy(C₁₋₄)alkyl, carboxyl,
 carboxyl(C₁₋₄)alkyl, amino (substituted with a substituent independently
 selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₄)alkyl
 (wherein amino is substituted with a substituent independently selected from the
 25 group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₄)alkyl,
 (halo)₁₋₃(C₁₋₄)alkoxy, hydroxy and hydroxy(C₁₋₄)alkyl; and, wherein
 heterocyclyl is optionally substituted with oxo); and, wherein R₉ is selected
 from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy(C₁₋₄)alkyl,
 carboxyl(C₁₋₄)alkyl, amino(C₁₋₄)alkyl (wherein amino is substituted with a
 30 substituent independently selected from the group consisting of hydrogen and
 C₁₋₄alkyl), hydroxy(C₁₋₄)alkyl, heterocyclyl(C₁₋₄)alkyl, aryl(C₁₋₄)alkyl and
 heteroaryl(C₁₋₄)alkyl (wherein the foregoing heterocyclyl, aryl and heteroaryl
 substituents are optionally substituted with one to four substituents

independently selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkoxy(C₁₋₄)alkyl, carboxyl, carboxyl(C₁₋₄)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₄)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₄)alkyl, (halo)₁₋₃(C₁₋₄)alkoxy, hydroxy and hydroxy(C₁₋₄)alkyl; and, wherein heterocyclyl is optionally substituted with oxo)).

6. The compound of claim 1 wherein R₂ is selected from the group consisting of -C₁₋₈alkyl- (optionally substituted with one to three substituents independently selected from the group consisting of halogen, hydroxy and oxo); aryl, heteroaryl, -(O-(CH₂)₁₋₆)₀₋₅-O-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-S-(CH₂)₁₋₆-O- and -NR₆- (wherein R₆, R₇ and R₈ are independently selected from the group consisting of hydrogen, C₁₋₄alkyl and C₁₋₄alkoxy(C₁₋₄)alkyl);

with the proviso that, if A and E are selected from a hydrogen substituted carbon atom, then R₂ is selected from the group consisting of -(O-(CH₂)₁₋₆)₁₋₅-O-, -(O-(CH₂)₁₋₆)₁₋₅-NR₆-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O- and -NR₆-(CH₂)₁₋₆-NR₇ -(CH₂)₁₋₆-NR₈- (wherein R₆, R₇ and R₈ are independently selected from the group consisting of hydrogen, C₁₋₄alkyl and hydroxy(C₁₋₄)alkyl).

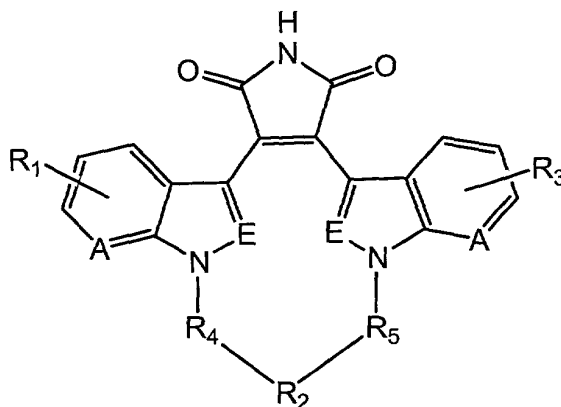
7. The compound of claim 1 wherein R₂ is selected from the group consisting of -C₁₋₈alkyl- (optionally substituted with one to two substituents independently selected from the group consisting of halogen, hydroxy and oxo); phenyl, pyridinyl, -(O-(CH₂)₂)₁₋₄-O-, -O-(CH₂)₂-NR₆-(CH₂)₂-O-, -O-(CH₂)₂-S-(CH₂)₂-O- and -NR₆- (wherein R₆, R₇ and R₈ are independently selected from the group consisting of hydrogen, C₁₋₃alkyl and C₁₋₂alkoxy(C₁₋₂)alkyl);

with the proviso that, if A and E are selected from a hydrogen substituted carbon atom, then R₂ is selected from the group consisting of -(O-(CH₂)₂)₁₋₄-O-, -(O-(CH₂)₂)₂-NR₆-, -O-(CH₂)₂-NR₆-(CH₂)₂-O- and

-NR₆-(CH₂)₂-NR₇-(CH₂)₂-NR₈- (wherein R₆, R₇ and R₈ are independently selected from the group consisting of hydrogen, C₁₋₃alkyl and hydroxy(C₁₋₂)alkyl).

8. The compound of claim 1 wherein R₁ and R₃ are independently selected from the group consisting of hydrogen, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl (wherein alkyl, alkenyl and alkynyl are optionally substituted with a substituent selected from the group consisting of C₁₋₄alkoxy, alkoxy(C₁₋₄)alkyl, carboxyl, carboxyl(C₁₋₄)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₄)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), (halo)₁₋₃, (halo)₁₋₃(C₁₋₄)alkyl, (halo)₁₋₃(C₁₋₄)alkoxy, hydroxy, hydroxy(C₁₋₄)alkyl and oxo), C₁₋₄alkoxy, C₁₋₄alkoxycarbonyl, (halo)₁₋₃(C₁₋₄)alkoxy, C₁₋₄alkylthio, aryl, heteroaryl (wherein aryl and heteroaryl are optionally substituted with a substituent selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, alkoxy(C₁₋₄)alkyl, carboxyl, carboxyl(C₁₋₄)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₄)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₄)alkyl, (halo)₁₋₃(C₁₋₄)alkoxy, hydroxy and hydroxy(C₁₋₄)alkyl), amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), cyano, halogen, hydroxy and nitro.
9. The compound of claim 1 wherein R₁ and R₃ are independently selected from the group consisting of hydrogen, C₁₋₄alkyl (optionally substituted with a substituent selected from the group consisting of C₁₋₄alkoxy, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), (halo)₁₋₃, hydroxy and oxo), C₁₋₄alkoxy, C₁₋₄alkoxycarbonyl, (halo)₁₋₃(C₁₋₄)alkoxy, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, hydroxy and nitro.

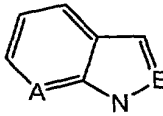
10. The compound of claim 1 wherein R_1 and R_3 are hydrogen.
11. The compound of claim 1 wherein a compound of Formula (I) is selected from a compound of Formula (Iaa):



Formula (Iaa)

wherein

- 5 A and E are independently selected from the group consisting of a hydrogen substituted

carbon atom and a nitrogen atom; wherein  is independently selected from the group consisting of 1H-indole, 1H-pyrrolo[2,3-b]pyridine and 1H-indazole;

- R_4 and R_5 are independently selected from C_{1-8} alkyl, C_{2-8} alkenyl and C_{2-8} alkynyl
10 optionally substituted with oxo;

- R_2 is selected from the group consisting of $-C_{1-8}$ alkyl-, $-C_{2-8}$ alkenyl-, $-C_{2-8}$ alkynyl-,
-O-(C_{1-8})alkyl-O-, -O-(C_{2-8})alkenyl-O-, -O-(C_{2-8})alkynyl-O-,
-C(O)-(C_{1-8})alkyl-C(O)- (wherein any of the foregoing alkyl, alkenyl and alkynyl
15 linking groups are straight carbon chains optionally substituted with one to four
substituents independently selected from the group consisting of C_{1-8} alkyl,
 C_{1-8} alkoxy, C_{1-8} alkoxy(C_{1-8})alkyl, carboxyl, carboxyl(C_{1-8})alkyl,
-C(O)O-(C_{1-8})alkyl, - C_{1-8} alkyl-C(O)O-(C_{1-8})alkyl, amino (substituted with a
substituent independently selected from the group consisting of hydrogen and

C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy, hydroxy(C₁₋₈)alkyl and oxo; and, wherein any of the foregoing alkyl, alkenyl and alkynyl linking groups
 5 are optionally substituted with one to two substituents independently selected from the group consisting of heterocyclyl, aryl, heteroaryl, heterocyclyl(C₁₋₈)alkyl, aryl(C₁₋₈)alkyl, heteroaryl(C₁₋₈)alkyl, spirocycloalkyl and spiroheterocyclyl (wherein any of the foregoing cycloalkyl, heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently
 10 selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl,
 15 (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein any of the foregoing heterocyclyl substituents are optionally substituted with oxo)), cycloalkyl, heterocyclyl, aryl, heteroaryl (wherein cycloalkyl, heterocyclyl, aryl and heteroaryl are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl,
 20 carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein heterocyclyl is
 25 optionally substituted with oxo), -(O-(CH₂)₁₋₆)₀₋₅-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -(O-(CH₂)₁₋₆)₀₋₅-NR₆-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-NR₆-, -(O-(CH₂)₁₋₆)₀₋₅-S-, -O-(CH₂)₁₋₆-S-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-S-, -NR₆-, -NR₆-NR₇-, -NR₆-(CH₂)₁₋₆-NR₇-, -NR₆-(CH₂)₁₋₆-NR₇-(CH₂)₁₋₆-NR₈-, -NR₆-C(O)-, -C(O)-NR₆-,
 30 -C(O)-(CH₂)₀₋₆-NR₆-(CH₂)₀₋₆-C(O)-, -NR₆-(CH₂)₀₋₆-C(O)-(CH₂)₁₋₆-C(O)-(CH₂)₀₋₆-NR₇-, -NR₆-C(O)-NR₇-, -NR₆-C(NR₇)-NR₈-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S-, -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S-, -NR₆-(CH₂)₁₋₆-S-(CH₂)₁₋₆-NR₇- and -SO₂- (wherein

R₆, R₇ and R₈ are independently selected from the group consisting of hydrogen, C₁₋₈alkyl, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl(C₁₋₈)alkyl, amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), hydroxy(C₁₋₈)alkyl, heterocyclyl(C₁₋₈)alkyl, aryl(C₁₋₈)alkyl and heteroaryl(C₁₋₈)alkyl (wherein the foregoing heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein heterocyclyl is optionally substituted with oxo));

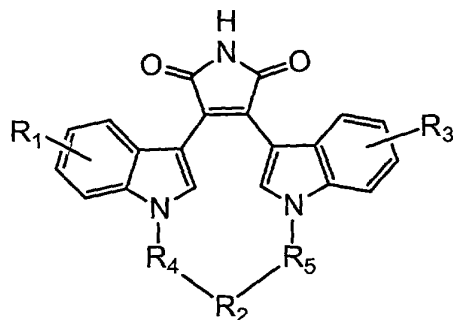
with the proviso that, if A and E are selected from a hydrogen substituted carbon atom, then R₂ is selected from the group consisting of -C₂₋₈alkynyl-, -O-(C₁₋₈)alkyl-O-, -O-(C₂₋₈)alkenyl-O-, -O-(C₂₋₈)alkynyl-O-, -C(O)-(C₁₋₈)alkyl-C(O)- (wherein any of the foregoing alkyl, alkenyl and alkynyl linking groups are straight carbon chains optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, -C(O)O-(C₁₋₈)alkyl, -C₁₋₈alkyl-C(O)O-(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy, hydroxy(C₁₋₈)alkyl and oxo; and, wherein any of the foregoing alkyl, alkenyl and alkynyl linking groups are optionally substituted with one to two substituents independently selected from the group consisting of heterocyclyl, aryl, heteroaryl, heterocyclyl(C₁₋₈)alkyl, aryl(C₁₋₈)alkyl, heteroaryl(C₁₋₈)alkyl, spirocycloalkyl and spiroheterocyclyl (wherein any of the foregoing cycloalkyl, heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and

C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein any of the foregoing heterocyclyl substituents are optionally substituted with oxo)), cycloalkyl (wherein cycloalkyl is optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl), -(O-(CH₂)₁₋₆)₁₋₅-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -(O-(CH₂)₁₋₆)₁₋₅-NR₆-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-NR₆-, -(O-(CH₂)₁₋₆)₀₋₅-S-, -O-(CH₂)₁₋₆-S-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-S-, -NR₆-NR₇-, -NR₆-(CH₂)₁₋₆-NR₇-, -NR₆-(CH₂)₁₋₆-NR₇-(CH₂)₁₋₆-NR₈-, -NR₉-C(O)-, -C(O)-NR₉-, -C(O)-(CH₂)₀₋₆-NR₆-(CH₂)₀₋₆-C(O)-, -NR₆-(CH₂)₀₋₆-C(O)-(CH₂)₁₋₆-C(O)-(CH₂)₀₋₆-NR₇-, -NR₆-C(O)-NR₇-, -NR₆-C(NR₇)-NR₈-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S-, -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S- and -NR₆-(CH₂)₁₋₆-S-(CH₂)₁₋₆-NR₇- (wherein R₆, R₇ and R₈ are independently selected from the group consisting of hydrogen, C₁₋₈alkyl, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl(C₁₋₈)alkyl, amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), hydroxy(C₁₋₈)alkyl, heterocyclyl(C₁₋₈)alkyl, aryl(C₁₋₈)alkyl and heteroaryl(C₁₋₈)alkyl (wherein the foregoing heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein heterocyclyl is optionally substituted with oxo); and, wherein R₉ is selected from the group

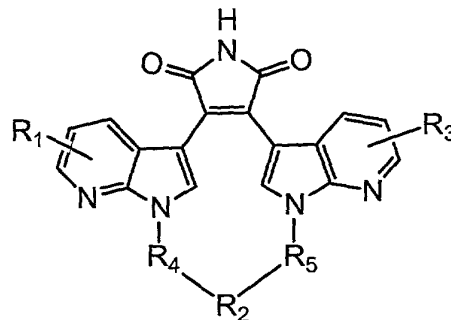
- consisting of C₁₋₈alkyl, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl(C₁₋₈)alkyl, amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), hydroxy(C₁₋₈)alkyl, heterocyclyl(C₁₋₈)alkyl, aryl(C₁₋₈)alkyl and heteroaryl(C₁₋₈)alkyl (wherein the foregoing heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein heterocyclyl is optionally substituted with oxo)); and,
- 15 R₁ and R₃ are independently selected from the group consisting of hydrogen, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl (wherein alkyl, alkenyl and alkynyl are optionally substituted with a substituent selected from the group consisting of C₁₋₈alkoxy, alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), (halo)₁₋₃, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy, hydroxy(C₁₋₈)alkyl and oxo), C₁₋₈alkoxy, C₁₋₈alkoxycarbonyl, (halo)₁₋₃(C₁₋₈)alkoxy, C₁₋₈alkylthio, aryl, heteroaryl (wherein aryl and heteroaryl are optionally substituted with a substituent selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl), amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), cyano, halogen, hydroxy and nitro;

and pharmaceutically acceptable salts thereof.

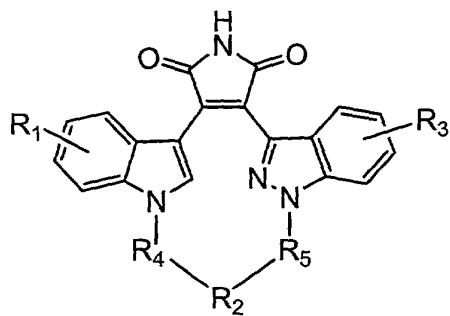
12. The compound of claim 1 wherein a compound of Formula (I) is selected from the group consisting of:



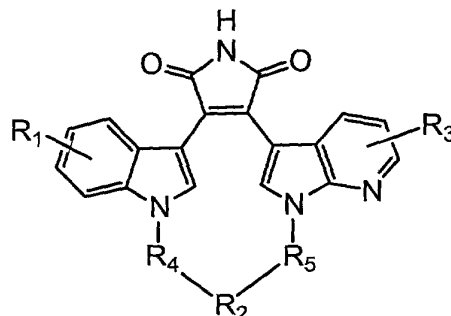
Formula (Ia);



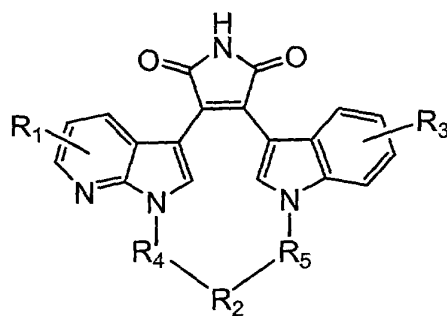
Formula (Ib);



Formula (If);



Formula (Ii); and,



Formula (Ij);

5 wherein

R_4 and R_5 are independently selected from C_{1-8} alkyl, C_{2-8} alkenyl and C_{2-8} alkynyl optionally substituted with oxo;

R_2 is selected from the group consisting of $-C_{1-8}$ alkyl-, $-C_{2-8}$ alkenyl-, $-C_{2-8}$ alkynyl-,
 10 $-O-(C_{1-8})$ alkyl-O-, $-O-(C_{2-8})$ alkenyl-O-, $-O-(C_{2-8})$ alkynyl-O-,

-C(O)-(C₁₋₈)alkyl-C(O)- (wherein any of the foregoing alkyl, alkenyl and alkynyl
 linking groups are straight carbon chains optionally substituted with one to four
 substituents independently selected from the group consisting of C₁₋₈alkyl,
 C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl,
 5 -C(O)O-(C₁₋₈)alkyl, -C₁₋₈alkyl-C(O)O-(C₁₋₈)alkyl, amino (substituted with a
 substituent independently selected from the group consisting of hydrogen and
 C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent
 independently selected from the group consisting of hydrogen and C₁₋₄alkyl),
 halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy, hydroxy(C₁₋₈)alkyl and
 10 oxo; and, wherein any of the foregoing alkyl, alkenyl and alkynyl linking groups
 are optionally substituted with one to two substituents independently selected from
 the group consisting of heterocyclyl, aryl, heteroaryl, heterocyclyl(C₁₋₈)alkyl,
 aryl(C₁₋₈)alkyl, heteroaryl(C₁₋₈)alkyl, spirocycloalkyl and spiroheterocyclyl
 (wherein any of the foregoing cycloalkyl, heterocyclyl, aryl and heteroaryl
 15 substituents are optionally substituted with one to four substituents independently
 selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl,
 carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently
 selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl
 (wherein amino is substituted with a substituent independently selected from the
 20 group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl,
 (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein any of the
 foregoing heterocyclyl substituents are optionally substituted with oxo)),
 cycloalkyl, heterocyclyl, aryl, heteroaryl (wherein cycloalkyl, heterocyclyl, aryl and
 heteroaryl are optionally substituted with one to four substituents independently
 25 selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl,
 carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently
 selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl
 (wherein amino is substituted with a substituent independently selected from the
 group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl,
 30 (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein heterocyclyl is
 optionally substituted with oxo), -(O-(CH₂)₁₋₆)₀₋₅-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-,
 -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -(O-(CH₂)₁₋₆)₀₋₅-NR₆-,
 -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-NR₆-, -(O-(CH₂)₁₋₆)₀₋₅-S-,

- $-O-(CH_2)_{1-6}-S-(CH_2)_{1-6}-O-$, $-O-(CH_2)_{1-6}-O-(CH_2)_{1-6}-S-$, $-NR_6-$, $-NR_6-NR_7-$,
 $-NR_6-(CH_2)_{1-6}-NR_7-$, $-NR_6-(CH_2)_{1-6}-NR_7-(CH_2)_{1-6}-NR_8-$, $-NR_6-C(O)-$, $-C(O)-NR_6-$,
 $-C(O)-(CH_2)_{0-6}-NR_6-(CH_2)_{0-6}-C(O)-$,
 $-NR_6-(CH_2)_{0-6}-C(O)-(CH_2)_{1-6}-C(O)-(CH_2)_{0-6}-NR_7-$, $-NR_6-C(O)-NR_7-$,
5 $-NR_6-C(NR_7)-NR_8-$, $-O-(CH_2)_{1-6}-NR_6-(CH_2)_{1-6}-S-$, $-S-(CH_2)_{1-6}-NR_6-(CH_2)_{1-6}-O-$,
 $-S-(CH_2)_{1-6}-NR_6-(CH_2)_{1-6}-S-$, $-NR_6-(CH_2)_{1-6}-S-(CH_2)_{1-6}-NR_7-$ and $-SO_2-$ (wherein
 R_6 , R_7 and R_8 are independently selected from the group consisting of hydrogen,
 C_{1-8} alkyl, C_{1-8} alkoxy(C_{1-8})alkyl, carboxyl(C_{1-8})alkyl, amino(C_{1-8})alkyl (wherein
amino is substituted with a substituent independently selected from the group
10 consisting of hydrogen and C_{1-4} alkyl), hydroxy(C_{1-8})alkyl, heterocyclyl(C_{1-8})alkyl,
aryl(C_{1-8})alkyl and heteroaryl(C_{1-8})alkyl (wherein the foregoing heterocyclyl, aryl
and heteroaryl substituents are optionally substituted with one to four substituents
independently selected from the group consisting of C_{1-8} alkyl, C_{1-8} alkoxy,
 C_{1-8} alkoxy(C_{1-8})alkyl, carboxyl, carboxyl(C_{1-8})alkyl, amino (substituted with a
15 substituent independently selected from the group consisting of hydrogen and
 C_{1-4} alkyl), amino(C_{1-8})alkyl (wherein amino is substituted with a substituent
independently selected from the group consisting of hydrogen and C_{1-4} alkyl),
halogen, (halo) $_{1-3}$ (C_{1-8})alkyl, (halo) $_{1-3}$ (C_{1-8})alkoxy, hydroxy and hydroxy(C_{1-8})alkyl;
and, wherein heterocyclyl is optionally substituted with oxo));
20 with the proviso that, if A and E are selected from a hydrogen substituted carbon atom,
then R_2 is selected from the group consisting of $-C_{2-8}$ alkynyl-, $-O-(C_{1-8})$ alkyl-O-,
 $-O-(C_{2-8})$ alkenyl-O-, $-O-(C_{2-8})$ alkynyl-O-, $-C(O)-(C_{1-8})$ alkyl-C(O)- (wherein any of
the foregoing alkyl, alkenyl and alkynyl linking groups are straight carbon chains
optionally substituted with one to four substituents independently selected from the
25 group consisting of C_{1-8} alkyl, C_{1-8} alkoxy, C_{1-8} alkoxy(C_{1-8})alkyl, carboxyl,
carboxyl(C_{1-8})alkyl, $-C(O)O-(C_{1-8})$ alkyl, $-C_{1-8}$ alkyl-C(O)O-(C_{1-8})alkyl, amino
(substituted with a substituent independently selected from the group consisting of
hydrogen and C_{1-4} alkyl), amino(C_{1-8})alkyl (wherein amino is substituted with a
substituent independently selected from the group consisting of hydrogen and
30 C_{1-4} alkyl), halogen, (halo) $_{1-3}$ (C_{1-8})alkyl, (halo) $_{1-3}$ (C_{1-8})alkoxy, hydroxy,
hydroxy(C_{1-8})alkyl and oxo; and, wherein any of the foregoing alkyl, alkenyl and
alkynyl linking groups are optionally substituted with one to two substituents
independently selected from the group consisting of heterocyclyl, aryl, heteroaryl,

heterocyclyl(C₁₋₈)alkyl, aryl(C₁₋₈)alkyl, heteroaryl(C₁₋₈)alkyl, spirocycloalkyl and spiroheterocyclyl (wherein any of the foregoing cycloalkyl, heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein any of the foregoing heterocyclyl substituents are optionally substituted with oxo)), cycloalkyl (wherein cycloalkyl is optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl), -(O-(CH₂)₁₋₆)₁₋₅-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-O-, -(O-(CH₂)₁₋₆)₁₋₅-NR₆-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-NR₆-, -(O-(CH₂)₁₋₆)₀₋₅-S-, -O-(CH₂)₁₋₆-S-(CH₂)₁₋₆-O-, -O-(CH₂)₁₋₆-O-(CH₂)₁₋₆-S-, -NR₆-NR₇-, -NR₆-(CH₂)₁₋₆-NR₇-, -NR₆-(CH₂)₁₋₆-NR₇-(CH₂)₁₋₆-NR₈-, -NR₉-C(O)-, -C(O)-NR₉-, -C(O)-(CH₂)₀₋₆-NR₆-(CH₂)₀₋₆-C(O)-, -NR₆-(CH₂)₀₋₆-C(O)-(CH₂)₁₋₆-C(O)-(CH₂)₀₋₆-NR₇-, -NR₆-C(O)-NR₇-, -NR₆-C(NR₇)-NR₈-, -O-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S-, -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-O-, -S-(CH₂)₁₋₆-NR₆-(CH₂)₁₋₆-S- and -NR₆-(CH₂)₁₋₆-S-(CH₂)₁₋₆-NR₇- (wherein R₆, R₇ and R₈ are independently selected from the group consisting of hydrogen, C₁₋₈alkyl, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl(C₁₋₈)alkyl, amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), hydroxy(C₁₋₈)alkyl, heterocyclyl(C₁₋₈)alkyl, aryl(C₁₋₈)alkyl and heteroaryl(C₁₋₈)alkyl (wherein the foregoing heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl,

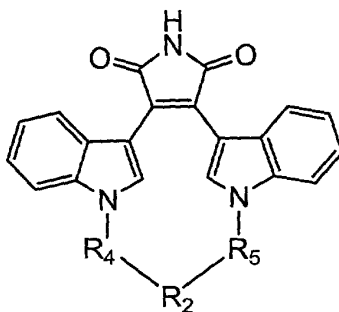
carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein heterocyclyl is optionally substituted with oxo); and, wherein R₉ is selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl(C₁₋₈)alkyl, amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), hydroxy(C₁₋₈)alkyl, heterocyclyl(C₁₋₈)alkyl, aryl(C₁₋₈)alkyl and heteroaryl(C₁₋₈)alkyl (wherein the foregoing heterocyclyl, aryl and heteroaryl substituents are optionally substituted with one to four substituents independently selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl; and, wherein heterocyclyl is optionally substituted with oxo)); and,

R₁ and R₃ are independently selected from the group consisting of hydrogen, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl (wherein alkyl, alkenyl and alkynyl are optionally substituted with a substituent selected from the group consisting of C₁₋₈alkoxy, alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), (halo)₁₋₃, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy, hydroxy(C₁₋₈)alkyl and oxo), C₁₋₈alkoxy, C₁₋₈alkoxycarbonyl, (halo)₁₋₃(C₁₋₈)alkoxy, C₁₋₈alkylthio, aryl, heteroaryl (wherein aryl and heteroaryl are optionally substituted with a substituent selected from the group consisting of C₁₋₈alkyl, C₁₋₈alkoxy, alkoxy(C₁₋₈)alkyl, carboxyl, carboxyl(C₁₋₈)alkyl, amino (substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), amino(C₁₋₈)alkyl

- (wherein amino is substituted with a substituent independently selected from the group consisting of hydrogen and C₁₋₄alkyl), halogen, (halo)₁₋₃(C₁₋₈)alkyl, (halo)₁₋₃(C₁₋₈)alkoxy, hydroxy and hydroxy(C₁₋₈)alkyl), amino (substituted with a substituent independently selected from the group consisting of hydrogen and
- 5 C₁₋₄alkyl), cyano, halogen, hydroxy and nitro;

and pharmaceutically acceptable salts thereof.

13. A compound of Formula (Ia1):

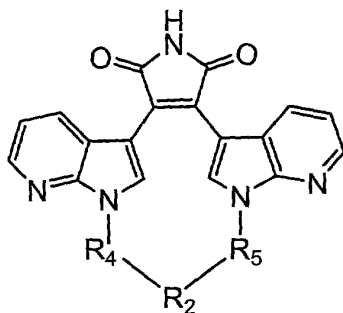


Formula (Ia1)

- 10 wherein R₄, R₂ and R₅ are dependently selected from:

R ₄	R ₂	R ₅
-(CH ₂) ₂ -	-O-(CH ₂) ₂ -O-	-(CH ₂) ₂ -;
-(CH ₂) ₂ -	-O-(CH ₂) ₂ -O-(CH ₂) ₂ -O-	-(CH ₂) ₂ -;
-(CH ₂) ₂ -	-O-(CH ₂) ₂ -O-(CH ₂) ₂ -O-(CH ₂) ₂ -O-	-(CH ₂) ₂ -;
-(CH ₂) ₂ -	-O-(CH ₂) ₂ -O-(CH ₂) ₂ -O-(CH ₂) ₂ -O-(CH ₂) ₂ -O-	-(CH ₂) ₂ -;
-(CH ₂) ₂ -	-O-(CH ₂) ₂ -N(Et)-(CH ₂) ₂ -O-	-(CH ₂) ₂ -;
-(CH ₂) ₂ -	-O-(CH ₂) ₂ -N(Me)-(CH ₂) ₂ -O-	-(CH ₂) ₂ -;
-(CH ₂) ₂ -	-O-(CH ₂) ₂ -N(<i>i</i> -Pr)-(CH ₂) ₂ -O-	-(CH ₂) ₂ -;
-(CH ₂) ₂ -	-N(Me)-(CH ₂) ₂ -N(Me)-(CH ₂) ₂ -N(Me)-	-(CH ₂) ₂ -;
-(CH ₂) ₂ -	-O-(CH ₂) ₂ -N(2-hydroxy-Et)-(CH ₂) ₂ -O-	-(CH ₂) ₂ -;
and,		
-(CH ₂) ₂ -	-O-(CH ₂) ₂ -O-(CH ₂) ₂ -N(Me)-	-(CH ₂) ₃ -.

14. A compound of Formula (Ib1):

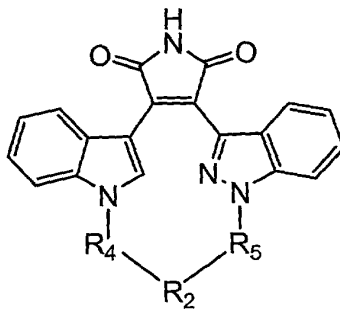


Formula (Ib1)

wherein R_4 , R_2 and R_5 are dependently selected from:

R_4	R_2	R_5
$-(CH_2)_2-$	$-O-(CH_2)_2-O-(CH_2)_2-O-$	$-(CH_2)_2-$;
$-(CH_2)_2-$	$-O-(CH_2)_2-O-(CH_2)_2-O-(CH_2)_2-O-$	$-(CH_2)_2-$;
$-(CH_2)_2-$	$-O-(CH_2)_2-O-(CH_2)_2-O-(CH_2)_2-O-(CH_2)_2-O-$	$-(CH_2)_2-$;
$-(CH_2)_2-$	$-O-(CH_2)_2-N(Et)-(CH_2)_2-O-$	$-(CH_2)_2-$;
$-(CH_2)_2-$	$-O-(CH_2)_2-S-(CH_2)_2-O-$	$-(CH_2)_2-$;
$-(CH_2)_5-$	$-NH-$	$-(CH_2)_5-$;
$-(CH_2)_5-$	$-N(Et)-$	$-(CH_2)_5-$;
$-(CH_2)_5-$	$-NH-$	$-(CH_2)_4-$;
$-(CH_2)_5-$	$-N(Et)-$	$-(CH_2)_4-$;
$-(CH_2)_4-$	$-2,6\text{-pyridinyl-}$	$-(CH_2)_4-$;
$-(CH_2)_4-$	$-C(O)-(CH_2)_2-$	$-(CH_2)_4-$;
$-(CH_2)_4-$	$-C(O)-$	$-(CH_2)_4-$;
$-CH_2-$	$-CH[R](OH)-(CH_2)_6-CH[R](OH)-$	$-CH_2-$;
and,		
$-(CH_2)_2-$	$-O-(CH_2)_2-O-$	$-(CH_2)_2-$.

15. A compound of Formula (If1):

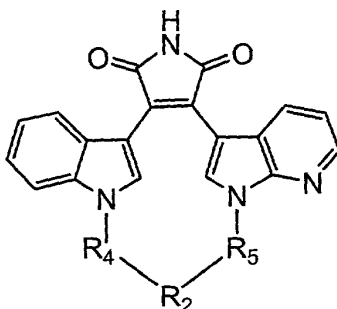


Formula (If1)

wherein R_4 , R_2 and R_5 are dependently selected from:

R_4	R_2	R_5
$-(CH_2)_2-$	$-O-(CH_2)_2-N(Me)-(CH_2)_2-O-$	$-(CH_2)_2-$;
$-(CH_2)_2-$	$-O-(CH_2)_2-N(Et)-(CH_2)_2-O-$	$-(CH_2)_2-$;
and,		
$-(CH_2)_2-$	$-O-(CH_2)_2-N(2-OMe-Et)-(CH_2)_2-O-$	$-(CH_2)_2-$.

16. A compound of Formula (Ii1):



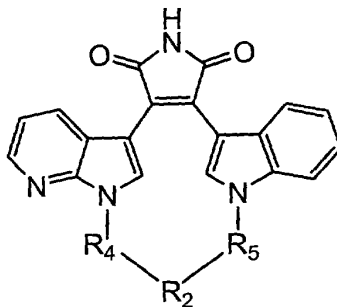
Formula (Ii1)

wherein R_4 , R_2 and R_5 are dependently selected from:

R_4	R_2	R_5
$-CH_2-$	-1,3-phenyl-	$-CH_2-$;
and,		
$-CH_2-$	-2,6-pyridinyl-	$-CH_2-$.

5

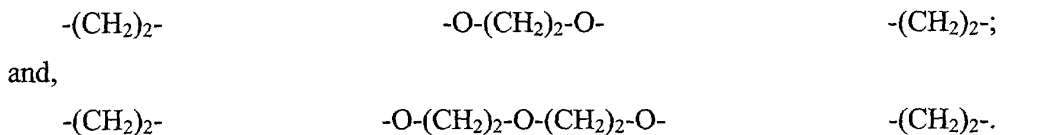
17. A compound of Formula (Ij1):



Formula (Ij1)

wherein R_4 , R_2 and R_5 are dependently selected from:

R_4	R_2	R_5
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18.
A pharmaceutical composition comprising a compound of claim 1 and a pharmaceutically acceptable carrier.
- 5
19.
A pharmaceutical composition made by mixing a compound of claim 1 and a pharmaceutically acceptable carrier.
20.
A method for preparing a pharmaceutical composition comprising mixing a compound of claim 1 and a pharmaceutically acceptable carrier.
- 10
21.
A method for treating or ameliorating a kinase mediated disorder comprising administering to a subject in need thereof a therapeutically effective amount of a compound of claim 1.
- 15
22.
The method of claim 21 wherein the disorder is mediated by selective inhibition of a kinase selected from the group consisting of protein kinase C and glycogen synthase kinase-3.
- 20
23.
The method of claim 22 wherein the kinase is selected from the group consisting of protein kinase C α , protein kinase C β -II, protein kinase C γ and glycogen synthase kinase-3 β .
- 25
24.
The method of claim 21 wherein the disorder is mediated by dual inhibition of at least two kinases selected from the group consisting of protein kinase C and glycogen synthase kinase-3.
- 30
25.
The method of claim 24 wherein at least two kinases are selected from the group consisting of protein kinase C α , protein kinase C β -II, protein kinase C γ and glycogen synthase kinase-3 β .

26. The method of claim 21 wherein the therapeutically effective amount of the compound of claim 1 is from about 0.001 mg/kg/day to about 300 mg/kg/day.
27. The method of claim 21 wherein the kinase mediated disorder is selected from the group consisting of cardiovascular diseases, diabetes, diabetes-associated disorders, inflammatory diseases, immunological disorders, dermatological disorders, oncological disorders and CNS disorders.
28. The method of claim 27 wherein cardiovascular diseases are selected from the group consisting of acute stroke, heart failure, cardiovascular ischemia, thrombosis, atherosclerosis, hypertension, restenosis, retinopathy of prematurity and age-related macular degeneration.
29. The method of claim 27 wherein diabetes is selected from the group consisting of insulin dependent diabetes and Type II non-insulin dependent diabetes mellitus.
30. The method of claim 27 wherein diabetes-associated disorders are selected from the group consisting of impaired glucose tolerance, diabetic retinopathy, proliferative retinopathy, retinal vein occlusion, macular edema, cardiomyopathy, nephropathy and neuropathy.
31. The method of claim 27 wherein inflammatory diseases are selected from the group consisting of vascular permeability, inflammation, asthma, rheumatoid arthritis and osteoarthritis.
32. The method of claim 27 wherein immunological disorders are selected from the group consisting of transplant tissue rejection, HIV-1 and PKC modulated immunological disorders.
33. The method of claim 27 wherein dermatological disorders are selected from the group consisting of psoriasis, hair loss and baldness.

34. The method of claim 27 wherein oncological disorders are selected from the group consisting of cancer, tumor growth, uncontrolled cell proliferation, proliferative angiopathy and angiogenesis.
- 5 35. The method of claim 27 wherein central nervous system disorders are selected from the group consisting of chronic pain, neuropathic pain, epilepsy, chronic neurodegenerative conditions, dementia, Alzheimer's disease, mood disorders, schizophrenia, manic depression and neurotraumatic, cognitive decline and ischemia-related diseases.
- 10 36. The method of claim 21 further comprising a method for use for a compound of claim 1 as an adjunct to chemotherapy and radiation therapy.
- 15 37. The method of claim 21 further comprising administering to a subject in need thereof a therapeutically effective amount of a pharmaceutical composition of claim 18.
- 20 38. The method of claim 37 wherein the therapeutically effective amount of a pharmaceutical composition of claim 18 is from about 0.001 mg/kg/day to about 300 mg/kg/day.
- 25 39. The method of claim 35 wherein ischemia-related diseases are those resulting from head trauma or transient ischemic stroke.